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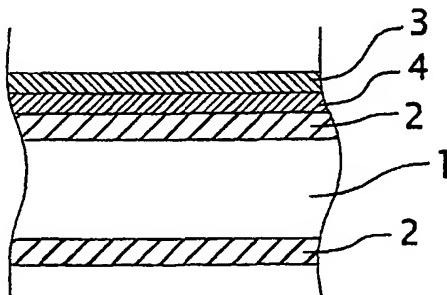
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(54) Title: BRAZING SHEET PRODUCT AND METHOD OF ITS MANUFACTURE



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(57) Abstract: A brazing sheet product and a method of manufacturing a brazing sheet product in which a layer comprising nickel is plated onto a surface of a clad layer made of an aluminium-silicon alloy containing silicon in the range of 2 to 18 weight %, wherein the surface is pre-treated by application of a bonding layer comprising zinc or tin. The application of the bonding layer may be by a zincate or a stannate treatment. The use of lead to promote wetting during brazing can be reduced or avoided, or other elements such as bismuth can be used.

**BRAZING SHEET PRODUCT AND METHOD OF ITS MANUFACTURE****FIELD OF THE INVENTION**

The invention relates to a method of manufacturing  
5 a brazing sheet product in which a layer comprising  
nickel is plated onto a surface of a clad layer made of  
an Al-Si alloy containing Si in the range of 2 to 18  
weight %. The invention also relates to a brazing sheet  
product obtained by the method and to a brazed assembly  
10 comprising at least one component made of the brazing  
sheet product.

**DESCRIPTION OF THE RELATED ART**

For the purpose of this invention brazing sheet is  
to be understood as a core sheet, for example of  
15 aluminium or aluminium alloy, having on at least one side  
a brazeable aluminium alloy. Typical brazeable aluminium  
alloys useful as such a clad layer are the Aluminium  
Association (AA) 4xxx-series alloys, typically having Si  
in the range of 2 to 18 weight %. The brazeable  
20 aluminium alloys may be coupled to the core alloy in  
various ways known in the art, for example by means of  
roll bonding, cladding, or semi-continuous or continuous  
casting.

Controlled Atmosphere Brazeing (CAB) and Vacuum  
25 Brazeing (VB) are the two main processes used for  
industrial scale aluminium brazeing. Industrial vacuum  
brazeing has been used since the 1950's, while CAB became

popular in the early 1980's after the introduction of the Nocoloc (trade mark) brazing flux. Vacuum brazing is an essentially discontinuous process and puts high demands on material cleanliness. The disruption of the oxide layer present is mainly caused by the evaporation of magnesium from the clad alloy. There is always more magnesium present in the furnace than necessary. The excess magnesium condenses on the cold spots in the furnace and has to be removed frequently. The capital investment for suitable equipment is relatively high.

CAB requires an additional process step prior to brazing as compared to VB, since a brazing flux has to be applied prior to brazing. CAB is essentially a continuous process in which, if the proper brazing flux is being used, high volumes of brazed assemblies can be manufactured. The brazing flux dissolves the oxide layer at brazing temperature allowing the clad alloy to flow properly. When the Nocoloc flux is used the surface needs to be cleaned thoroughly prior to flux application. To obtain good brazing results the brazing flux has to be applied on the total surface of the brazed assembly. This can cause difficulties with certain types of assemblies because of their design. For example, because evaporator type heat exchangers have a large internal surface, problems can arise because of poor access to the interior. For good brazing results the flux has to adhere to the aluminium surface before brazing.

Unfortunately the brazing flux after drying can easily fall off due to small mechanical vibrations. During the brazing cycle, corrosive fumes such as HF are generated. This puts a high demand on the corrosion resistance of 5 the materials applied for the furnace.

Ideally, a material should be available that can be used for CAB but does not have the requirements and defects of the brazing flux application. Such a material can be supplied to a manufacturer of brazed assemblies 10 and is ready to use directly after shaping of the assembly parts. No additional brazing fluxing operations have to be carried out. Presently, only one process for flux-less brazing is used on an industrial scale. The material for this process can be for example standard 15 brazing sheet made from an AA3xxx-series core alloy clad on both sides with a cladding of an AA4xxx-series alloy. Before the brazing sheet can be used the surface has to be modified in such a way that the naturally occurring oxide layer does not interfere during the brazing cycle. 20 The method of achieving good brazing is to deposit a specific amount of nickel on the surface of the clad alloy. If properly applied, the nickel reacts, presumably exothermically, with the underlying aluminium. The nickel can be applied by using a shim of nickel 25 between the two parts to be joined or can be deposited by electroplating. When electroplating is used the adherence of the nickel should be sufficient to withstand

typical shaping operations being used in for example heat exchanger manufacture.

- The processes for nickel-plating of aluminium brazing sheet are known from each of US-A-3,970,237, US-5 A-4,028,200, US-A-4,164,454, and SAE-paper no. 880446 by B.E. Cheadle and K.F. Dockus. According to these documents, nickel is preferably deposited in combination with lead. The lead addition is used to improve the wettability of the clad alloy during the brazing cycle.
- 10 An important characteristic of these plating processes is that the nickel is preferentially deposited on the silicon particles of the clad alloy. To obtain sufficient nickel for brazing on the surface, the clad alloy should contain a relatively large number of silicon
- 15 particles to act as nuclei for the nickel deposition. It is believed that to obtain sufficient nucleation sites before pickling a part of the aluminium in which the silicon particles are embedded should be removed by chemical and/or mechanical pre-treatment. This is
- 20 believed a necessary condition to obtain a sufficient nickel coverage to serve as nuclei for the wetting action of the brazing or clad alloy. On a microscopic scale the surface of the Si-containing cladding of the brazing sheet is covered with nickel globules.
- 25 Some other disclosures of Ni-plating found in the prior art literature will be mentioned below.

General textbook by Wernick and Pinner, "The Surface

Treatment and Finishing of Aluminium and its Alloys", 5th edition, Volume 2, pp.1023-1071. This textbook describes in general immersion processes for plating on aluminium.

Paper by the Bureau of Mines Technology, "Aluminium 5 Soft-Soldering", 2301 N.T.I.S. Tech Notes (manufacturing), 1985, January, No.1G, Springfield, VA, USA, pp.12-13.

This paper describes a method of manufacturing aluminium for soft-soldering wherein the aluminium surfaces are joined by conventional tin-lead solder. The method 10 includes firstly cleaning the aluminium surface carefully prior to the zinc application. Secondly a thin zinc coat is deposited and subsequently electroplated with an alloy of nickel-copper. After the nickel-copper plating has been accomplished, soldering using normal 15 procedures can be accomplished.

FR-A-2,617,868 describes a method of manufacturing aluminium product with a brazeable surface coating of tin or a tin-bismuth alloy, wherein the product is provided with an intermediate layer. This intermediate layer is 20 composed of a first layer of zinc and a second layer of nickel, which nickel has been deposited by electrolysis from a neutral electrolyte. Here, the underlying aluminium or aluminium alloy is not melted in the brazing process.

25 SUMMARY OF THE INVENTION

It is an object of the present invention to provide a method of manufacturing nickel-plated brazing sheet

product, comprising a core provided on at least one side with a clad of an Al-Si alloy comprising Si in the range of 2 to 18 weight %, by which method good adhesion of the surface of the clad layer to the nickel is obtained.

5 It is also an object of the invention to provide a brazing sheet product having a core sheet and filler material, to be melted during brazing, comprising a clad layer of an Al-Si alloy and a nickel layer on the clad layer, in which there is good adhesion of the nickel  
10 layer to the clad layer.

In accordance with the invention in one aspect there is provided a method of manufacturing a brazing sheet product, comprising the step of plating a layer comprising nickel onto a surface of a sheet comprising a core sheet and a clad layer on the core sheet, the clad layer being made of an aluminium alloy containing silicon in an amount in the range 2 to 18% by weight and said surface being a surface of the clad layer, the method including a pre-treatment of said surface before the  
15 plating step. This method is characterised in that the pre-treatment comprises the step of applying a bonding layer comprising zinc or tin on said surface.

By the zinc or tin pretreatment step of the invention an effective bond between the aluminium alloy  
20 clad layer and the layer comprising nickel is formed, the bond remaining effective during subsequent deformation of the brazing sheet, for example by bending. The coverage

of the nickel layer is no longer dependent on the surface characteristics of the bare clad layer. Furthermore the method may be carried out in a continuous process. The product obtained by this method is suitable for flux-less  
5 brazing under controlled atmosphere conditions.

The invention is based in part on the insight that to obtain a well-bonded nickel layer on the Si-containing clad layer of the brazing sheet product so that the bond remains effective under large deformation, pretreatment  
10 of the clad layer is extremely important. The prior art processes apparently aimed at applying the nickel in a distributed form, principally to the silicon particles at the surface of the clad layer, rather than trying to achieve a uniform nickel layer. In the present invention  
15 the surface of the Si-containing clad alloy is altered in such way that the nickel coverage is independent of the silicon particles at its surface. The nickel plating does not take place on the silicon particles but on the applied layer comprising zinc or tin. Since the nickel  
20 thus is deposited on the total surface of the clad layer the necessary reaction before brazing can take place much more easily as compared to the process of the prior art. The zinc or tin applied does not interfere at all during  
25 the brazing process, and may contain a component to assist the brazing, as described below. Since the nickel is deposited smoothly and uniformly on the surface, the use of lead to promote wetting during brazing can be

reduced or avoided, or other elements such as bismuth may be used for this purpose. A further important advantage of the nickel or nickel-lead deposited smoothly and uniformly on the surface is that the total amount of 5 nickel to be applied in order to achieve good flux-less brazing can be reduced. Another advantage is that the complete surface coverage avoids any difficulty caused by aluminium oxide at the surface of the clad layer.

While it is in general known to apply a zinc layer 10 prior to nickel-plating of articles, it is believed that this has not been done in a nickel-plated aluminium alloy clad brazing sheet product, in which as discussed above it has been thought necessary to plate the nickel directly on the Si-containing clad layer.

15 Very good results may be obtained with an immersion zincate treatment or immersion stannate treatment, often also referred to as displacement plating. A further advantage is that this treatment lends itself to application in a continuous process operation.

20 Preferably the duration of the zincate treatment or stannate treatment is in the range of 1 to 300 seconds.

25 Preferably the temperature of the bath during the zincate treatment or stannate treatment is in the range of 10 to 50°C, and more preferably in the range of 15 to 30°C.

Zincate treatments are known per se in the art for applying layers onto aluminium, for example as known from

"Oppervlaktebehandelingen van aluminium" by T. van der Klis and J.W. du Mortier published by the Vereniging voor Oppervlaktetchnieken voor Materialen, Bilthoven, NL, 3rd edition 1992, pp. 406-409. A simple basic composition  
5 for a zincate pickle comprises 40-50 g/l ZnO and 400-500 g/l NaOH. Also, other commercial available zincate baths can be used, for example ChemTec (tradename) 024202, also known as the Bondal process, and ChemTec (tradename) 024195, also known as a cyanide-free Bondal process.

10 Stannate treatments are known in the art for depositing a layer on aluminium to facilitate soldering, to improve electrical conductivity, and also to give a lubricated surface to aluminium alloy pistons for internal combustion engines during the running-in period.  
15 Typical alkaline stannate solutions comprise 5-300 g/l sodium or potassium stannate.

Preferably in the method to the invention the applied layer comprising zinc or tin has a thickness up to 0.5 $\mu$ m, more preferably up to 0.3 $\mu$ m (300nm), and most 20 preferably in the range of 0.01 to 0.15 $\mu$ m (10-150nm). In the best results obtained a thickness of about 30nm has been used. A coating thickness of greater than 0.5 $\mu$ m requires a prolonged treatment time, e.g. for displacement plating, and is thought to have no further 25 advantages for improving the adhesion.

The zinc or tin layer applied in the method of the invention may be essentially a pure zinc or tin layer or

may be primarily zinc or tin (e.g. at least 50 weight %). Minor amounts of impurity elements or deliberately added elements may be present, as discussed in more detail below. Typically impurity elements are present at less than 10%, more usually less than 5% by weight in the zinc or tin layer. The zinc or tin layer may contain less than 1 % of other elements.

Following the application of the bonding layer according to the method of the invention the aluminium brazing sheet is typically plated with nickel, nickel-lead, nickel-cobalt or nickel-lead-cobalt by electroplating in an alkaline solution. Good results may be obtained when the electroplating process for nickel or nickel-lead deposition comprises one or more of:

- 15 (a) bath temperature 20-70°C, preferably 20-30°C;
- (b) pH 7.0-12.0, preferably pH 10.0-12.0, and more preferably about 10.5;
- (c) current density of 0.1-10.0 A/dm<sup>2</sup>, preferably 0.5-4.0 A/dm<sup>2</sup>;
- 20 (d) plating time 1 to 300 s, preferably 30 to 100 s;
- (e) bath composition comprising 3-200 g/l nickel sulfate, preferably 50 g/l nickel sulfate, 10-100 g/l nickel chloride, preferably 50 g/l nickel chloride, 60-300 g/l sodium citrate, preferably 100 g/l sodium citrate, 0.05-10.0 g/l lead acetate, preferably 1.0 g/l lead acetate, 5-150 ml/l ammonium hydroxide (30% by weight), preferably 75

- ml/l ammonium hydroxide. As alternative for the sodium citrate 60-300 g/l sodium gluconate, preferably 150 g/l sodium gluconate may be used, for the lead acetate 0.05-5 g/l lead citrate or 5 bismuth lactate, preferably 1.0 g/l lead citrate or bismuth lactate may be used. In case of nickel-cobalt or nickel-lead-cobalt plating the bath composition may further comprise cobalt chloride in the range of 10-100 g/l, preferably 50 g/l.
- 10 Using these parameters in combination with the bonding layer in accordance with the invention, a well-bonded layer comprising essentially nickel or nickel-lead is applied to the brazing sheet, the bonding remaining effective under large deformation of the nickel-plated
- 15 brazing sheet and the deposition of the plating layer being independent of the silicon particles at the surface of the clad layer. A further advantage is that it is possible to perform a continuous process.
- Alternatively, after the application of the bonding 20 layer according to the method of the invention the aluminium brazing sheet is plated with nickel or nickel-lead by electroplating in an acidic solution. Good results may be obtained when in the electroplating 25 process for nickel or nickel-lead deposition the parameters comprise one or more of:-
- (a) bath temperature 20-70°C, preferably 40-60°C;

- (b) pH in the range of 3 to 5, preferably 4 to 5;
- (c) current density of 0.1-10.0 A/dm<sup>2</sup>, preferably 0.5 to 5.0 A/dm<sup>2</sup>;
- (d) plating time 1 to 300 s, preferably 20 to 100 s;
- 5 (e) bath composition comprising 5-400 g/l nickel sulphate, preferably 240-300 g/l nickel sulphate, 10-100 g/l nickel chloride, preferably 40-60 g/l nickel chloride, 5-100 g/l boric acid, preferably 25-40 g/l boric acid.
- 10 Such an electroplating process is often referred to in the act as the Watt's process. Using these parameters in combination with the bonding layer in accordance with the invention, a well-bonded layer comprising essentially nickel or nickel-lead may be applied to the brazing sheet, the bonding remaining effective under large deformation of the nickel-plated brazing sheet and the deposition of the plating layer being independent of the silicon particles at the surface of the clad layer. A further advantage is that it is possible to perform a
- 15 continuous process.

Alternatively, following the application of the bonding layer according to the method of the invention the aluminium brazing sheet is nickel or nickel-lead plated by electroplating in an acid solution comprising nickel or nickel-lead using alkylsulphonic acid electrolytes, and preferably methanesulphonic acid.

Alternatively, following the application of the

bonding layer according to the method of the invention the aluminium brazing sheet is plated with nickel or nickel-lead by electroplating in a sulfamate solution or in a lead sulfamate solution. Typically the sulfamate 5 solution comprises 50-500 g/l nickel sulfamate, 0.05-30 g/l lead sulfamate, 15-50 g/l boric acid, and optionally wetting agents. Bath temperatures are in the range of 20 to 70°C.

Alternatively, following the application of the 10 bonding layer according to the method of the invention the aluminium brazing sheet is plated with nickel or nickel-lead by electroplating in a fluoborate or in a lead fluoborate ( $Pb(BF_4)_2$ ) solution. Typically nickel fluoborate is present in the range 50-500 g/l, optionally 15 lead fluoborate in the range of 0.05-30.0 g/l, and further optionally fluoboric acid in the range 1-50 g/l, boric acid 15-50 g/l, and further optionally a wetting agent. Bath temperatures are in the range of 20 to 80°C, and preferably 40 to 70°C. An advantage is that this 20 solution, like some others here described, does not require the use of ammonium hydroxide.

Alternatively, following the application of the bonding layer according to the method of the invention the aluminium brazing sheet is plated with nickel or 25 nickel-lead by electroplating in a bath comprising 50-500 g/l nickel acetate, 0.05-30 g/l lead acetate, 15-50 g/l boric acid, up to 200 ml/l glycolic acid (70%), 20-100

g/l sodium acetate, and optionally wetting agents.

The invention further provides a brazed assembly comprising at least one component made of the brazing sheet product produced by the method in accordance with 5 the invention described above.

In another aspect of the invention there is provided a brazing sheet product having a core sheet, a clad layer on the core sheet made of an aluminium alloy containing silicon in an amount in the range 2 to 18% by 10 weight, and a layer comprising nickel on the outer surface of the clad layer, characterised by a layer comprising zinc or tin as a bonding layer between the outer surface of the clad layer and the layer comprising nickel.

15 The layer comprising nickel is preferably an electroplated layer. The adhesion of the layer comprising nickel applied on the layer comprising zinc or tin is excellent and can withstand relatively severe shaping operations without the occurrence of 20 delamination.

Preferably in the brazing sheet product according to the invention the layer comprising zinc or tin has a thickness up to 0.5 $\mu\text{m}$ , more preferably up to 0.3 $\mu\text{m}$ , and most preferably in the range of 0.01 to 0.15 $\mu\text{m}$ . A 25 coating thickness of greater than 0.5 $\mu\text{m}$  requires a prolonged treatment time for plating.

Preferably in this brazing sheet product the layer

comprising nickel has a thickness up to 2.0 $\mu\text{m}$ , preferably up to 1.0 $\mu\text{m}$ , and more preferably up to 0.5 $\mu\text{m}$ . A coating thickness of greater than 2.0 $\mu\text{m}$  requires a prolonged treatment time for plating, may result in wrinkling of 5 the nickel layer and is thought to have no further advantages during brazing. A preferred minimum thickness for this Ni-containing layer is 0.3 $\mu\text{m}$ .

Preferably in the brazing sheet product the material which on brazing becomes molten, known as filler 10 material, in particular the nickel layer and/or the zinc or tin layer comprise one or more elements to reduce the surface tension of the molten brazing alloy during brazing. In the invention it has been found surprisingly that contrary to the teaching of the prior art, it is not 15 necessary to add lead as an alloying element to the Ni-layer in order to promote the wetting action of the brazing alloy. Nevertheless, lead and other suitable elements, for which bismuth is preferred to most, may be added to the nickel layer or the zinc or tin layer or to 20 both. This has various advantages from the manufacturing point of view of the brazing sheet.

In the filler material as a whole therefore, there may be present, in weight %, at least one of

Bi 0.01 to 0.5, preferably 0.05 to 5  
25 Mg 0.2 to 2.0  
Sb 0.01 to 0.5, preferably 0.05 to 5.

The zinc or tin layer itself may thus comprise one or more additional elements selected from the group consisting of bismuth, lead, lithium and antimony. The amount of the additional element or elements in total may 5 be up to 50%, but preferably is less than 25%, e.g. in the range 1 to 25%.

The clad layer may comprise, in weight %, Si in the range of 2 to 18 %, preferably 7 to 18%, and Mg in the range of up to 6%. Preferably the Mg is in the range of 10 0.5 to 5%. Further alloying elements may be added such as, but not limited to, Cu, Zn, Sr and Mn in suitable ranges. It has been found that in use of the brazing sheet the presence of Mg in the cladding has no detrimental effects during brazing. This is a major 15 improvement over known brazing sheets. It allows for the design of a cladding which may contribute to the strength of the total brazing sheet product. Further, it allows that Mg-containing brazing sheet may be applied in both Vacuum Brazing and flux-less Controlled Atmosphere 20 Brazing. The latter possibility has many economical and technical advantages. The brazing sheet according to the invention may be readily used in the existing industrial brazing lines without the change of relevant process parameters, such as temperature and processing time.

25 In another embodiment the clad layer comprises, in weight %, Si in the range of 2 to 18 % and Zn in the range of up to 5%. Preferably the Zn is in the range of

0.5 to 3%. Further alloying elements may be added such as, but not limited to, Mg, Cu and Mn in suitable ranges.

In accordance with the invention it has been found that when this brazing sheet is used the presence of Zn in the 5 cladding has no detrimental effects during brazing. This is a major improvement over known brazing sheets. It allows for the design of a cladding which may contribute to the strength of the total brazing sheet product.

Further, the brazing sheet product wherein the cladding 10 contains Zn as a deliberate alloying element may be applied in both Vacuum Brazing and flux-less Controlled Atmosphere Brazing, both processing being used on an industrial scale.

Typically in the brazing sheet product according to 15 the invention the core sheet is an aluminium alloy but the invention is not limited to this and any suitable material may be used. It is necessary that the core sheet has a melting point higher than that of the filler material (i.e. all layers which melt during brazing to 20 generate the brazing alloy).

In one preferred embodiment, the core sheet is an aluminium alloy comprising Mg in a range of up to 8%. In a preferred embodiment Mg is in a range of 0.5 to 5.0 wt.%. Further alloying elements may be added such as, 25 but not limited to, Cu, Zn, Bi, V, Fe, Zr, Ag, Si, Ni, Co and Mn in suitable ranges. It has been found that when the brazing sheet of the invention is used, the presence

of Mg in the clad layer has no detrimental effects during brazing. This is a major improvement over the known brazing sheets. The diffusion of Mg from the core to the cladding during the manufacturing of the brazing sheet 5 product itself and its application in a subsequent brazing process, appears to have no detrimental effects on the brazeability of the brazing sheet in accordance with the invention. This allows for the design of high strength brazing sheet product having an aluminium core 10 sheet having Mg in the given range as a strengthening element. The product may be applied in both Vacuum Brazing and flux-less Controlled Atmosphere Brazing, both processes being used on an industrial scale.

In the brazing sheet product according to the 15 invention the core sheet may be coupled to the clad layer via an intermediate layer. The benefits of having such an intermediate layer or interlayer are described in for example US-A-2,821,014, the contents of which are incorporated here by reference.

20 In a further aspect of the invention there is provided in a method a manufacturing a brazed assembly using brazing sheet product in accordance with the invention, comprising the steps of:-

(a) shaping parts of which at least one is made from 25 the brazing sheet product of the invention as set out above;

- (b) assembling the parts into the assembly;
  - (c) brazing the assembly under a vacuum or in an inert atmosphere in the absence of a brazing-flux at elevated temperature for a period long enough for
    - 5 melting and spreading of the cladding alloy;
  - (d) cooling the brazed assembly. The cooling rate may be in the range of typical brazing furnace cooling rates. Typical cooling rates are a cooling rate of at least 10°C/min or more.
- 10 In dependence on the material, particularly aluminium alloy, of the core sheet the process may include the further processing step (e) of ageing of the brazed and cooled assembly in order to optimise the mechanical and corrosion properties of the assembly.
- 15 BRIEF INTRODUCTION OF THE DRAWINGS

The invention will now be illustrated by several non-limitative examples, and with reference to the drawings, wherein:

Fig. 1 is a schematic longitudinal section showing  
20 the structure of brazing sheet product according to the state of the art;

Fig. 2 is a schematic longitudinal section showing the structure of brazing sheet product according to the invention;

25 Fig. 3 is a schematic longitudinal section showing the structure of brazing sheet product according to the invention;

Figs. 4A and 4B are SEM photographs of the surface after Ni-plating according to the prior art and in accordance with the present invention respectively.

#### DESCRIPTION OF PREFERRED EMBODIMENTS

5 Fig. 1 shows schematically brazing sheet in accordance with the prior art as would be obtained by the process in accordance with for example US-A-3,970,237. The brazing sheet product consists of a core sheet 1 clad on one or both sides clad with a cladding layer 2  
10 comprising an Al-Si-alloy. On top of the clad 2 a thin nickel layer 3, preferably a nickel-lead layer, is applied by means of electroplating.

Fig. 2 shows schematically brazing sheet in accordance with the present invention where between the 15 cladding layer 2 and the Ni-layer 3 a further layer 4 of zinc or tin is applied, the advantages of which are set out above. In Fig. 2 the layers 4 and 3 have been shown on only one side of the brazing sheet, but it will be immediately apparent to the skilled person that they may 20 also be applied on both sides of the brazing sheet product. The composition of the various layers and their advantages have been set out above.

Fig. 3 shows schematically a further brazing sheet in accordance with the present invention which has the 25 layers of Fig. 2 and a further intermediate layer 5 between the core sheet 1 and the cladding layer 2 on both sides. In Fig. 3 the layers 4 and 3 have been shown on

only one side of the brazing sheet, but it will be immediately apparent to the skilled person that they may also be applied on both sides of the brazing sheet product. In addition the intermediate layer 5 may also 5 be applied on one side of the brazing sheet, preferably on the side comprising also the layers 4 and 3. The possible compositions of the various layers and their advantages have been set out above.

Fig. 4A is a SEM back-scatter photograph 10 (magnification 1000x) of the surface of 0.5µm Ni-plated brazing sheet in accordance with the prior art as described in US-A-3,970,237. Globules of Ni can be observed which are preferentially deposited on the Si-particles of the clad layer of the brazing sheet. Fig. 15 4B represents a SEM back-scatter photograph (magnification 1000x) of the surface of 0.5µm Ni-plated brazing sheet in accordance with the invention having a Zn layer under the Ni layer and here it can be observed that the Ni is deposited much more uniformly and smoothly 20 over the entire surface giving rise to several advantages during brazing as set out above.

#### EXAMPLES

##### Example 1

25 On a laboratory scale of testing was carried out on aluminium brazing sheet manufactured from an AA3003 core alloy clad on both sides with an AA4045-series clad

alloy, having a total thickness of 0.5 mm, the thickness of each clad being 10.9% of the total thickness. The composition of these alloys is given in Table 1.

5 Table 1 Weight %

	AA 3003	AA 4045
Si	0.6 max.	9.0 - 11.0
Fe	0.7 max.	0.8 max.
Cu	0.05 - 0.20	0.30 max.
Mn	1.0 - 0.20	0.05 max.
Mg	-	0.05 max.
Zn	0.10 max.	0.10 max.
Ti	-	0.20 max.
impurities	each 0.05 total 0.15	each 0.05 total 0.15
balance	aluminium	aluminium

Each sample was treated by the following sequential process steps (see also Table 2),

- cleaning by immersion for 180 s in ChemTec 30014 (a  
10 commercial available bath),  
rinsing,  
alkaline etching for 20 s in ChemTec 30203 (a  
commercial available bath component),  
rinsing,  
15 optionally desmutting for 4 s in an acidic  
oxidising solution, typically 25-50 vol.% nitric acid,

comprising ChemTec 11093 (a commercial available pickle activator) at ambient temperature, followed by rinsing, optional zincate immersion using ChemTec 024202 for 12 s at room temperature, followed by rinsing,  
5 nickel electroplating, and rinsing.

For the nickel electroplating two different types of solutions were used, a basic bath and an acid bath, see also Table 2.

10 The acid bath comprised 270 g/l nickel sulphate, 50 g/l nickel chloride, 30 g/l boric acid. The plating conditions at 50°C were such that a nickel layer of 2.0 $\mu\text{m}$  is present after the plating process using a current density of 5 A/dm<sup>2</sup>. This acid bath is also known as the  
15 Watt's process.

The basic bath comprised 50 g/l nickel sulphate, 50 g/l nickel chloride, 100 g/l sodium citrate, 1 g/l lead acetate, and 75 ml/l ammonium hydroxide (30%). The plating conditions at 26°C were such that a plating time  
20 of 50 s resulted in a nickel-lead plated layer of 0.5 $\mu\text{m}$  thickness using a current density of 3 A/dm<sup>2</sup>, and a plating time of 200 s resulted in a nickel-lead plated layer of 2.0 $\mu\text{m}$  thickness.

25 The nickel plated specimens have been tested for adhesion using the Erichsen dome test (5 mm), and the T-

bend test. A value assessment is then given to the adhesion where: (-) = poor, ( $\pm$ ) = fair, and (+) = good. The morphology of the nickel layer applied has been investigated using SEM/EDX, where: U = uniform nickel 5 layer (shiny appearance), and (G) = nickel globules preferentially deposited on the silicon particles (dull appearance). Further the brazeability has been assessed.

On a laboratory scale of testing the brazing tests were carried out in a small quartz furnace. Small coupons of 10 25 mm x 25 mm were cut from the nickel-plated sheets. A small strip of an AA3003 alloy measuring 30 mm x 7 mm x 1 mm was bent in the centre to an angle of 45° and laid on the coupons. The strip on the coupon samples were heated under flowing nitrogen, with heating from room 15 temperature to 580°C, dwell time at 580°C for 1 minute, cooling from 580°C to room temperature. The brazing process was judged on possible formation of wrinkles, capillary depression and fillet formation. An overall assessment was given where: (-) = poor brazeability, (-/±) = fair brazeability, ( $\pm$ ) = good brazeability, and (+) = excellent brazeability. The results obtained are 20 summarised in Table 2.

From the results of Table 2 it can been seen that in case of a zinc immersion pretreatment step, a uniform 25 nickel or nickel-lead layer is obtained having a shiny

appearance. Further it can be seen that with a basic bath a zinc immersion pretreatment is required to obtain a good adhesion of the electroplated nickel layer.

Further it can be seen that a nickel-lead layer of 0.5 $\mu\text{m}$

5 has better brazeability characteristics than a layer of 2.0 $\mu\text{m}$  thickness; in the latter case wrinkles have been observed. The brazeability of the material obtained via the basic bath route has better brazeability characteristics (but still acceptable) than material

10 obtained via the acid bath route, possibly due to the presence of lead in the electroplated layer.

Table 2 Experimental conditions and results.

Plating bath	acid	acid	acid	basic	basic	basic	basic
Desmutting [s]	4	4	-	4	-	4	-
Zinc immersion time [s]	-	12	12	-	12	12	12
Nickel plating time [s]	120	120	120	200	200	50	50
Adhesion	-	+	+	-	+	+	+
Brazeability	-/±	-/±	-/±	±	±	+	+
Morphology	G	U	U	G	U	U	U

#### Example 2

On a laboratory scale of testing aluminium brazing sheets manufactured from an AA3003 core alloy clad on one side with AA4000-series clad alloys of six different

compositions (see Table 3), and having a total thickness of 0.5 mm, and a clad thickness of about 50 $\mu$ m, were treated as set out in Table 4.

The treatment consisted of:-

5       cleaning by immersion for 180 s in ChemTec 30014 (a commercial available bath),

rinsing,

alkaline etching for 20 s in ChemTec 30203 (a commercial available pickle activator) at ambient 10 temperature, rinsing,

optionally desmutting for 4 s in an acidic oxidizing bath, typically 25-50 vol% nitric acid, comprising ChemTec 11093 at ambient temperature, followed by rinsing,

15       optional zincate immersion using ChemTec 024202 for 12 s at room temperature followed by rinsing,

nickel electroplating, and rinsing.

For the nickel electroplating two different types 20 of solutions have been used: a basic bath with lead and a basic bath without lead.

The lead-containing basic bath comprised 50 g/l nickel sulphate, 50 g/l nickel chloride, 30 g/l sodium citrate, 1 g/l lead-acetate and 75 ml/l ammonium 25 hydroxide (30%). The plating conditions at 26°C were such

that a plating time of 50 s resulted in a nickel-lead plated layer of 0.5 $\mu\text{m}$  thickness using a current density of 3 A/dm<sup>2</sup>. The process using this bath is in Table 4 referred to as "L+".

5       The lead-free basic bath comprised 50 g/l nickel sulphate, 50 g/l nickel chloride, 30 g/l sodium citrate, and 75 ml/l ammonium hydroxide (30%). The plating conditions at 26°C were such that a plating time of 50 s resulted in a nickel-lead plated layer of 0.5 $\mu\text{m}$  thickness  
10 using a current density of 3 A/dm<sup>2</sup>. The process using this bath is in Table 4 referred to as "L-".

The samples were tested as in Example 1 set out above. The results are set out in Table 4. In this table "n.t." stands for not tested.

15       From the results in Table 4 it can be seen that a zincate treatment in accordance with the invention succeeds in achieving good adhesion of the Ni-plated layer. From the results of clad alloy no. 1 it can be seen that omitting the desmutting step still results in a  
20 good adhesion of the Ni-plated layer. An excellent brazeability can be achieved using a wide variety of cladding compositions. The presence of Mg in clad layer no. 3 has no detrimental effect of the brazeability of the brazing sheet. The presence of Zn in clad layer no.  
25 5 has no detrimental effect of the brazeability of the

brazing sheet.

Table 3 Composition of the clad alloy, in weight %.

Balance Al and inevitable impurities.

Alloy	Si	Fe	Cu	Mn	Mg	Zn	Ti	Bi	Sr
1	10.0	0.3	<0.01	<0.02	<0.02	<0.02	0.003	-	-
2	10.0	0.3	<0.01	<0.02	<0.02	<0.02	0.003	-	-
3	9.6	0.25	<0.01	<0.02	1.32	<0.02	0.003	-	-
4	7.6	0.3	<0.01	<0.02	<0.02	<0.02	0.003	-	0.02
5	7.6	0.35	<0.01	<0.02	<0.02	1.02	0.003	-	-
6	12.4	0.2	<0.01	0.06	<0.02	<0.02	0.003	-	-

Table 4 Pretreatment applied and testing results.

Clad alloy no.	Clean.	Etching	Desmut.	Zincate	Ni-plating	Adhesion	Braze-ability
1	yes	yes	yes	yes	L-	+	-
1	yes	yes	no	yes	L-	+	-
2	yes	yes	yes	yes	L+	+	+
2	yes	yes	yes	no	L+	-	n.t.
3	yes	yes	yes	yes	L+	+	+
3	yes	yes	yes	no	L+	-	+
4	yes	yes	yes	yes	L+	+	+
4	yes	yes	yes	no	L+	-	n.t.
5	yes	yes	yes	yes	L+	+	+
5	yes	yes	yes	no	L+	-	+
6	yes	yes	yes	yes	L+	+	+
6	yes	yes	yes	no	L+	-	n.t.

Example 3

5        In this example, bismuth was included in the zinc bonding layer, in order to reduce the surface tension of the molten brazing alloy formed by the clad layer and the layers above it during brazing. The sheet product on which the bonding layer was applied consisted of a core  
10      of alloy AA3003 (78.2% by weight) and two clad layers of alloy AA4045 (10.9% by weight each), total thickness 0.5 mm.

The displacement plating baths used in several trials were as shown in Table 5, which also shows the

results obtained in the adhesion test and brazeability tests carried out as in Example 1. The process steps were:-

- a. cleaning of clad surface, using bath of 35 g/l ChemTec 30014, 50°C, 180 s
- b. thorough rinsing
- c. alkaline etching, using bath of 30 g/l ChemTech 30203, 50°C, 20 s
- d. thorough rinsing
- e. desmutting, using 50% HNO<sub>3</sub>, room temperature, 60 s
- f. thorough rinsing
- g. zincate immersion, see Table 5, to apply Zn-Bi layer of about 35µm (about 0.25 g/m<sup>2</sup>)
- h. nickel plating, using the lead-free basic bath described above in Example 2, 26°C, 3 A/dm<sup>2</sup>, 50 s, corresponding to Ni layer thickness of 0.5µm (4.5 g/m<sup>2</sup>) .

Table 5

Zn-Bi bath	1	2	3	4
Sodium hydroxide g/l	120	120	120	120
Zinc oxide g/l	20	20	20	20
Potassium sodium tartrate g/l	50	50	50	50
Sodium nitrate g/l	1	1	1	1
Bismuth (III) oxide g/l	0	10	2	1
Plating time s	12	12	12	12
Appearance of 'bonding' layer	silver metallic	black powdery	grey	silver metallic
Composition of ZnBi layer from ICP data g/m <sup>2</sup>		Bi: 0.7 Zn: 0.35		Bi: 0.05 Zn: 0.25
Nickel plating	50 s	50 s	50 s	50 s
Adhesion test	excellent	very poor	poor	excellent
Brazeability test	-	+	+	+

The composition of the layer from bath 3 was not  
 5 measured. ICP stands for induced coupled plasma.

The results here show that the inclusion in the  
 thin bonding layer of even a small amount of Bi, as an  
 example of an element reducing surface tension of the  
 molten brazing alloy, achieves a significant effect on  
 10 brazing performance. However the presence of Bi in the  
 bonding layer reduces the adhesion effect achieved, while  
 still permitting brazing. The weight of Zn in the layer  
 should preferably be at least 50%, more preferably at  
 least 75%. Clearly the amount of addition of an element

such as Bi which can be achieved in the thin bonding layer is small, but the effect is significant, and a lead-free product was obtained. Avoidance of lead has environmental benefits.

5

Example 4

This example shows how, following the application of a zinc bonding layer as in Examples 1 and 2, an electroplated nickel layer containing Bi, but no Pb, may 10 be applied, resulting in a product having good adhesion of the nickel layer and good brazeability. Bismuth is not easily maintained in a stable nickel-plating solution, without sludge formation.

The nickel plating bath used has the composition of  
15 Table 6:

Table 6

Compound	g/l
Nickel sulphate	142
Ammonium sulphate	34
Nickel chloride	30
Sodium citrate	140
Sodium gluconate	30
Bismuth ions	1

Following the procedure of Examples 1 and 2, using  
 a product having a clad of AA4045 alloy, with a Zn  
 5 bonding layer, electroplating of a Ni-Bi layer was  
 performed at 57°C according to Table 7:-

Table 7

Current Density A/dm <sup>2</sup>	time s	Ni g m <sup>-2</sup> (ICP)	Bi g m <sup>-2</sup> (ICP)	adhesion	brazeability
3	50	9.1	0.66	+	+
6	25	10.4	0.50	+	+
10	15	9.5	0.46	+	+

- 10 The current efficiency of Ni deposition appears to be  
 100%.

The samples plated at 3 A/dm<sup>2</sup> showed some black spots, but the samples plated at the higher current densities have excellent appearance. About 0.5 g m<sup>-2</sup> bismuth was deposited. The bismuth content of the 5 deposited alloy layer can easily be varied, e.g. by lowering the bismuth concentration in the plating bath, to give a lower Bi content.

This plating bath has many advantages compared to the standard Pb-containing baths:-

- 10 no ammonia fumes
- more practical operating temperatures, typically 40 to 70°C
- high current density
- bismuth can easily be replenished to the plating
- 15 bath.

Further, standard chemicals were employed.

Having now fully described the invention, it will be apparent to one of ordinary skill in the art that many changes and modifications can be made without departing 20 from the spirit or scope of the invention as herein described.

CLAIMS

1. A method of manufacturing a brazing sheet product, comprising the step of plating a layer comprising nickel onto a surface of a sheet comprising a core sheet and a clad layer on the core sheet, the clad layer being made of an aluminium alloy containing silicon in an amount in the range 2 to 18% by weight and said surface being a surface of the clad layer, the method including a pre-treatment of said surface before the plating step,  
characterised in that the pre-treatment comprises the step of applying a bonding layer comprising zinc or tin on said surface.
2. A method according to claim 1, wherein said bonding layer is applied by a zincate treatment or a stannate treatment.
3. A method according to claim 2, wherein said bonding layer is applied by an immersion zincate treatment or an immersion stannate treatment.
- 20 4. A method according to claim 2 or 3, wherein the duration of the zincate or stannate treatment is in the range 1 to 300 s.
5. A method according to any one of claims 2 to 4, wherein the temperature of the zincate or stannate treatment is in the range of 10 to 50°C.

6. A method according to any one of claims 1 to 5, wherein said bonding layer has a thickness of not more than 1 $\mu$ m, preferably not more than 0.3 $\mu$ m.

7. A method according to claim 6, wherein said bonding layer has a thickness in the range 10 to 150nm.

8. A method according to any one of claims 1 to 7, wherein said layer comprising nickel is applied by electroplating.

9. A method according to claim 8, wherein said electroplating is carried out in a sulfamate solution.

10. A method according to claim 9, wherein said sulfamate solution contains lead, whereby said nickel layer contains lead.

11. A method according to any one of claims 8 to 10, wherein said layer comprising nickel is applied by electroplating in which one or more process parameters are selected from:-

- (a) electroplating bath temperature 20-70°C, preferably 20-30°C;
- 15 (b) electroplating bath pH 7.0-12.0, preferably pH 10.0-12.0, and more preferably about 10.5;
- (c) current density of 0.1-10.0 A/dm<sup>2</sup>, preferably 0.5-4.0 A/dm<sup>2</sup>;
- (d) plating time 1 to 300 s, preferably 30 to 100 s;
- 25 (e) bath composition comprising 3-200 g/l nickel

sulfate, 10-100 g/l nickel chloride, 60-300 g/l sodium citrate, preferably 100 g/l sodium citrate, 0.05-10.0 g/l lead acetate, and 5-150 ml/l ammonium hydroxide (calculated as 30% ammonium hydroxide  
5 solution).

12. A method according to claim 8, wherein said layer comprising nickel is applied by electroplating in which one or more process parameters are selected from:-

- (a) electroplating bath temperature 20-70°C, preferably 10 20-30°C;
- (b) electroplating bath pH in the range of 3 to 5, preferably 3 to 4, and more preferably about 3;
- (c) current density of 0.1-10.0 A/dm<sup>2</sup>, preferably 0.5 to 5.0 A/dm<sup>2</sup>;
- 15 (d) plating time 1 to 300 s, preferably 20 to 100 s;
- (e) bath composition comprising 5-400 g/l nickel sulphate, preferably 240-300 g/l nickel sulphate, 10-100 g/l nickel chloride, preferably 40-60 g/l nickel chloride, 5-100 g/l boric acid, preferably 20 25-40 g/l boric acid.

13. A method according to any one of claims 1 to 11, wherein the amount of silicon in said clad layer is in the range 7 to 18% by weight.

14. Brazing sheet product having a core sheet (1), a 25 clad layer (2) on said core sheet (1) made of an

aluminium alloy containing silicon in an amount in the range 2 to 18% by weight, and a layer (3) comprising nickel on the outer surface of said clad layer, characterised by a layer (4) comprising zinc or tin as a 5 bonding layer between said outer surface of said clad layer and said layer comprising nickel.

15. Brazing sheet product according to claim 14, wherein said clad layer (2) has discrete silicon-rich particles exposed at said outer surface thereof, and said 10 layer (3) comprising nickel is bonded both to said silicon-rich particles and to the areas of said outer surface between said silicon-rich particles, so as to form a continuous layer on said outer surface.

16. Brazing sheet product according to claim 14 or 15, 15 wherein said bonding layer (4) is an electroplated layer.

17. Brazing sheet product according to any one of claims 14 to 16, wherein said bonding layer (4) has a thickness of not more than 0.5 $\mu\text{m}$ , preferably not more than 0.3 $\mu\text{m}$ .

20 18. Brazing sheet product according to claim wherein said bonding layer (4) has a thickness in the range 20 to 150nm.

19. Brazing sheet product according to any one of claims 14 to 18, wherein said layer (3) comprising nickel 25 has a thickness of not more than 2.0 $\mu\text{m}$ , preferably not

more than 1.0 $\mu$ m.

20. Brazing sheet product according to any one of claims 14 to 19, wherein taken together said clad layer and all layers exterior thereto, have a composition containing at least one of the following elements,
- Bi in the range 0.01 to 0.5% by weight, preferably 0.05 to 0.5%
- Mg in the range 0.2 to 2.0% by weight
- Sb in the range 0.01 to 0.5% by weight, preferably 0.05 to 0.5%.

- 10 21. Brazing sheet product according to any one of claims 14 to 20, wherein said bonding layer (4) contains by weight an amount not more than 50%, and preferably not more than 25%, in total of one or more elements selected from bismuth, lead, lithium and antimony.

- 15 22. Brazing sheet product according to any one of claims 14 to 21, wherein said clad layer (2) contains by weight Mg in an amount in the range of up to 8%, preferably in the range of 0.5 to 5%.

- 20 23. Brazing sheet product according to any one of claims 14 to 22, wherein said clad layer (2) contains by weight Zn in an amount in the range of up to 5%, preferably in the range of 0.5 to 3%.

24. Brazing sheet product according to any one of 25 claims 14 to 23, wherein said core sheet (1) is an

aluminium alloy.

25. Brazing sheet product according to claim 24,  
wherein said core sheet (1) is an aluminium alloy  
comprising Mg in an amount in the range of up to 8%.

5 26. Brazing sheet product according to any one of  
claims 14 to 25, wherein said core sheet (1) is coupled  
to said clad layer (2) via an intermediate layer (5).

27. An assembly of components joined by brazing, at  
least one said components being a brazing sheet product  
10 according to any one of claims 14 to 26.

28. Method of manufacturing an assembly of brazed  
components, comprising the steps of:

- (a) forming said components of which at least one is  
made from brazing sheet product according to any  
15 one of claims 14 to 27;
- (b) assembling the components into an assembly;
- (c) brazing the assembly under a vacuum or in an inert  
atmosphere in the absence of a brazing-flux at  
elevated temperature for a period long enough for  
20 melting and spreading of the clad layer;
- (d) cooling the brazed assembly.

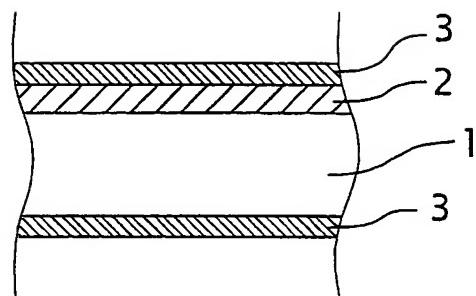


Fig. 1

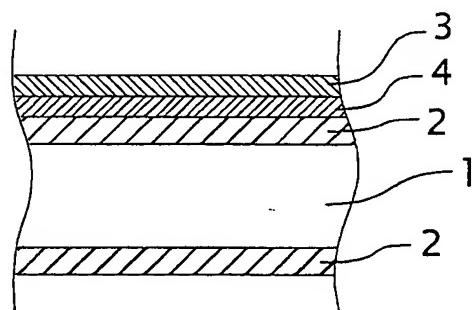


Fig. 2

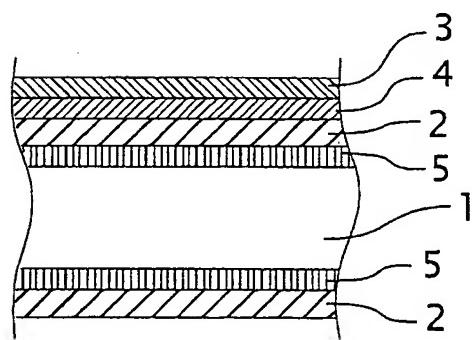


Fig. 3

2/2

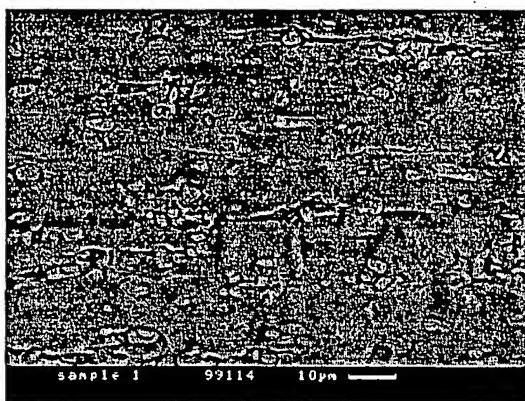


Fig. 4B

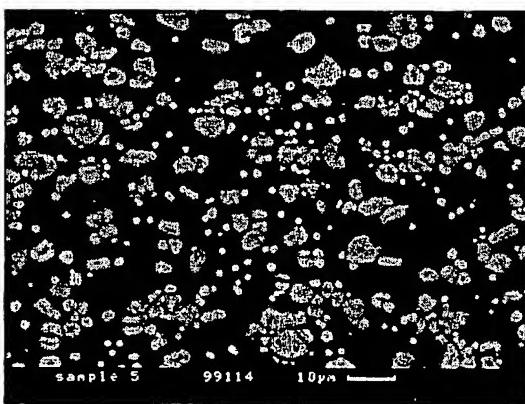


Fig. 4A